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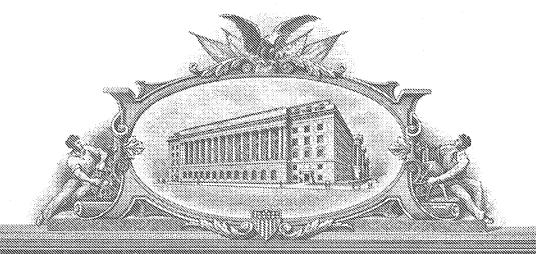
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United States Patent and Trademark Office

July 07, 2005

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APPLICATION NUMBER: 60/542,987 FILING DATE: February 09, 2004

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PROVISIONAL APPLICATION FOR PATENT COVER SHEET

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c).

100

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TITLE OF THE INVENTION (500 characters max)									
Immune Modulation through Targeting of the MINOR Gene									
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	ENCLOS	SED APPLICATION PAR	TS (check all	that apply)		<b>.</b>			
Specification Number of Pages  Drawing(s) Number of Sheets  Application Data Sheet. See 37 CFR 1.76				D(s), Numberther (specify)					
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT									
Applicant claims small entity status. See 37 CFR 1.27.  A check or money order is enclosed to cover the filing fees.						G FEE int (\$)			
The Director is herby authorized to charge filing fees or credit any overpayment to Deposit Account Number:					\$	80.00			
Payment by credit card. Form PTO-2038 is attached.									
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Respectfully submitted, SIGNATURE TYPED or PRINTED NAM	X Lather A	(Page 1 of bakalyar	D. Ri	ate <u>09-</u> EGISTRATION <i>appropriate</i> ) ocket Number	1 NO. <u>4</u>	-04 15,282 06			

TELEPHONE 410-516-8300

USE ONLY FOR FILING A PROVISIONAL APPLICATION FOR PATENT

This collection of information is required by 37 CFR 1.51. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 8 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop Provisional Application, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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[Page 2 of 2]

# CERTIFICATE OF EXPRESS MAILING EXPRESS MAILING LABEL NO.

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## Johns Hopkins University School of Medicine Office of Technology Licensing

## Report of Invention Disclosure Form

This form is to be completed and submitted to the JHU office of Licensing and Technology Development by anyone who believes they have developed a new invention. The purpose of this form is to enable OTL to evaluate whether legal protection to the invention will be sought and/or commercialization pursued. In order for this Report of Invention to be processed by LTD, it must be signed and dated by all inventors, and by the JHU Department Director(s) for all departments involved with the development of this invention. OTL can not process this report until it is complete with all necessary signatures found in Sections A, B and/or C. Visit the LTD web site at <a href="http://www.hopkinsmedicine.org/lbd/otl/RptInv.html">http://www.hopkinsmedicine.org/lbd/otl/RptInv.html</a> for HTML and Word 97 downloadable formats of this form.

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I	NVENTION INFORMATION		
Title of Invention:			
Immune Modulation through Targeting of	the MINOR gene	C OTT 1	
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Are you an HHMI employee or investiga	tor? Yes No		
Are you a KKI employee or investigator?	Yes No	onal Inventors se	ction for each
Additional inventors: X Yes	No If yes, please complete Additi	Onal miventors se	otion for outin
inventor.			
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JHU Ref.: 4406

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Page 2 JHU Ref.: 4406

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Are you a KKI emp	employee or investigat bloyee or investigator?	Yes No	.:				

## INVENTION DESCRIPTION

Describe the invention completely, using the outline given below.

## 1. Abstract of the Invention [Briefly describe the invention]

A number of vaccination strategies utilize dendritic cells (DCs) to immunize. While DCs are potent initiators of immune responses, their utility as vaccines may be limited by their relatively short in vivo lifespans. We have discovered a new gene in DCs that regulates DC apoptosis and have developed a strategy to inhibit expression of this gene, and by doing so, have shown that we can significantly enhance immune responses. This gene, termed , MINOR, for Mitogen Induced Nuclear Orphan Receptor, is a member of the Nur77 family of apoptosis-inducing genes, and its expression is highly and selectively upregulated in mature DCs, and, our data suggest that it plays a role in natural DC apoptosis. In order to enhance DC survival and function, we have developed a novel approach of inhibiting DC apoptosis via small interfering RNA (siRNA) technology. Our data suggest that inhibition of this gene leads to improvement of ex vivo DC vaccines and also utilize our system of bone marrow transplantation (BMT) with gene modified hematopoietic stem cells (HSCs) to analyze its effects on de novo generation of DCs in vivo.

2. Problem Solved [Describe the problem solved by this invention]

Dendritic cell vaccines have been developed for therapeutic use by generating the DCs both in vivo and in vitro, through various methods. However, these strategies have not been highly effective. Improving vaccination strategies for tumors is a significant goal of immunotherapy. It now appears that DC vaccines can elicit strong immune responses, but they are limited, in part by their short lifespans in vivo. While much emphasis has been placed in studying antigen (Ag) uptake, processing and presentation as well as costimulatory signal delivery by DCs, little is known about regulation of DC lifespan. Through investigating the unique pattern of gene expression, in DCs, we have identified one whose expression may be at least partly responsible for limiting the efficacy of DC vaccines due to its observed apoptosis-inducing effects. We show that by inhibiting this gene that we can prolong survival of DCs and also enhance immune responses. Thus, this is a novel approach to improving DC vaccines. (Claims on attached page)

3. Novelty [Identify those elements of the invention that are new when compared to the current state of the art] This report describes a new gene that is important for DC function. Expression of the new gene, MINOR, regulates apoptosis in DCs and is likely to be a limiting factor in immunogenicity of these cells. Inhibition of this gene provides a novel means to prolong the immune response in a number of settings. Potentiation of gene expression may also provide a means to inhibit the immune response in order to target hyper or autoimmune type processes.

## 4. Detailed Description of the invention:

On a separate page(s), attach a detailed description of how to make and use the invention. The description must contain sufficient detail so that one skilled in the same discipline could reproduce the invention. Include the following as necessary:

- 1- data pertaining to the invention;
- 2- drawings or photographs illustrating the invention;
- 3- structural formulae if a chemical;
- 4- procedural steps if a process
- 5- a description of any prototype or working model;

In general, a manuscript that has been prepared for submission to a journal will satisfy this requirement.

5. Workable Extent/Scope [Describe the future course of related work, and possible variations of the present invention in terms of the broadest scope expected to be operable; if a compound, describe substitutions, breadth of substituents, derivatives, salts etc., if DNA or other biological material, describe modifications that are expected to be operable, if a machine or device, describe operational parameters of the device or a component thereof, including alternative structures for performing the various functions of the machine or device]

Future plans will center on applying inhibition of this gene in order to enhance immunogenicity in a number of different types of vaccine therapies. In addition, investigation into immunosuppression by potentiating expression of this gene is another therapeutic goal.

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## 2. Problem Solved - continued:

#### Claims:

- 1. A new dendritic cell-selective gene, MINOR, capable of inducing apoptosis in dendritic cells
- 2. Manipulation of MINOR to develop vaccines for cancer through ex vivo DC-based vaccines using molecular targeting, e.g, siRNA.
- 3. Manipulation of MINOR to develop vaccines for cancer through ex vivo DC-based vaccines using molecular targeting, e.g, signal transduction inhibitors.
- 4. Manipulation of MINOR to develop vaccines for cancer through ex vivo DC-based vaccines using cell based targeting, e.g, cellular delivery of MINOR targeting molecules.
- 5. The use of MINOR reagents to classify and/or isolate specific populations of DCs.
- 6. Manipulation of MINOR to develop vaccines for cancer through blood or marrow transplantation with gene modified cells for transplant, by means of introduction of MINOR inhibiting sequences or genes.
- 7. Manipulation of MINOR to develop vaccines for cancer through blood or marrow transplantation with cells for transplant treated with MINOR-targeted signal transduction inhibitors or by using cell-based inhibition for delivery to the stem-progenitor cells used for transplant.
- 8. Manipulation of MINOR by molecular targeting, e.g, siRNA, to develop antigen specific immunity
- 9. Manipulation of MINOR by molecular targeting, e.g, signal transduction inhibitors, to develop antigen specific immunity.
- 10. Manipulation of MINOR by cell based targeting, e.g, cellular delivery of MINOR targeting molecules to develop antigen specific immunity.
- 11. Manipulation of MINOR to enhance general immunogenicity of cells by cell based targeting, e.g, cellular delivery of MINOR targeting molecules.
- 12. Manipulation of MINOR to enhance general immunogenicity of cells by gene based approaches for targeting MINOR in immune cells.
- 13. Manipulation of MINOR to develop vaccines for viral and or bacterial disease for therapy and prophylaxis.
- 14. Manipulation of MINOR to develop vaccines for immunodeficiencies of unknown origin through inhibition of MINOR to enhance immunogenicity, by siRNA, small molecular weight compounds, or cell based approaches.
- 15. Manipulation of MINOR in antigen presenting cells by gene or cell-based approaches, to develop novel adjuvants for vaccines for cancers and infectious disease.
- 16. Potentiation of MINOR as an immunosuppressant for autoimmune or hyperimmune syndromes or to induce tolerance

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